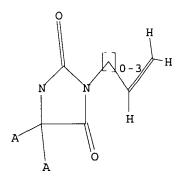
09/535,348 Page 1

=> d l1 L1 HAS NO ANSWERS STR L1



Structure attributes must be viewed using STN Express query preparation.

=> d his

(FILE 'HOME' ENTERED AT 10:57:45 ON 24 OCT 2003)

FILE 'REGISTRY' ENTERED AT 10:57:53 ON 24 OCT 2003

STRUCTURE UPLOADED L1

L20 S L1

27 S L1 FULL L3

27 S L3 AND CAPLUS/LC L4 L5

9 S L3 AND CAOLD/LC

FILE 'CAOLD' ENTERED AT 10:59:16 ON 24 OCT 2003

L6 2 S L5

FILE 'CAPLUS' ENTERED AT 10:59:57 ON 24 OCT 2003

L7 27 S L3 ANSWER 9 OF 27 CAPLUS COPYRIGHT 2003 ACS on STN

1984:571168 CAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER: 101:171168

Study of some N-3 substituted and N-1, N-3 TITLE:

disubstituted derivatives of cyclohexanospirohydantoin

Pedregal, Carmen; Trigo, Gregorio G.; Espada, Modesta; AUTHOR (S):

Elguero, Jose; Vincent, Emile Jean; Faure, Robert Dep. Quim. Org. Farm., Fac. Farm., Madrid, Spain

CORPORATE SOURCE: Journal of Heterocyclic Chemistry (1984), 21(2), SOURCE:

477-80

CODEN: JHTCAD; ISSN: 0022-152X

DOCUMENT TYPE: Journal French LANGUAGE:

CASREACT 101:171168 OTHER SOURCE(S):

GT

Alkylation of spirocyclohexanehydantoin I (R = R1 = H) gave 72-81% I (R = AB H; R1 = Me, Et, Bu, CH2CH:CH2, CH2Ph, CH2CH2NEt2). In the presence of phase transfer catalyst Bu4NBr, dialkylation occurred, to give 68-94% I (R = R1 = Me, Et, Bu, CH2CH:CH2, CH2Ph, CH2CH2NEt) using the same reagents. I (R = Bu, R1 = CH2Ph; R = CH2Ph, R1 = Bu) were prepd. in 93% yield by alkylation of I (R = H; R1 = Bu, CH2Ph) in presence of Bu4NBr. Treating I (R = R1 = H) with 0.5 equiv Br(CH2)3Br gave 73% of the di(spirocyclohexanehydantoinyl) compd II.

882-66-6P 92357-90-9P ΙT

RL: SPN (Synthetic preparation); PREP (Preparation) (prepn. of)

882-66-6 CAPLUS RN

1,3-Diazaspiro[4.5]decane-2,4-dione, 3-(2-propenyl)- (9CI) (CA INDEX CN NAME)

$$CH_2-CH$$